

## CLAIM AMENDMENTS

## 1. (canceled)

2. (currently amended) A DNazyme which binds to GATA-3 mRNA and functionally inactivates it, which comprises:

- a catalytic domain with the nucleotide sequence

~~GGCTAGCTA-CA-AC-GA~~ GGCTAGCTACAACGA SEQ ID NO: 154 or a modified sequence with comparable biological effect, which cleaves the GATA-3 mRNA at every purine:pyrimidine binding site to which it is bonded,

- a right substrate binding domain adjoining the 3' end of the catalytic domain and

- a left substrate binding domain adjoining the 5' end of the catalytic domain, both substrate binding domains being respectively complementary to two regions of the GATA 3 mRNA so that they hybridize with the mRNA, and

- ~~[[are]]~~ which is active in vivo.

3. (currently amended) A DNazyme according to claim 2, which comprises the sequence hgd 40 GTGGATGGA GGCTAGCTACAACGA  
GTCCTTGGAG GTCTTGGAG SEQ ID NO: 40.

1                   4. (previously presented) A DNzyme according to claim 2  
2    which cleaves the catalytic domain of the GATA-3 mRNA at every  
3    purine:uracil binding site.

1                   5. (previously presented) A DNzyme according to claim 2  
2    which is stabilized against decomposition within the organism by  
3    introduction of a 3'-3' inversion.

1                   6. (previously presented) A DNzyme according to claim 2  
2    which is stabilized against decomposition within the organism by  
3    introduction of modified nucleotides or nucleotide compounds.

1                   7. (previously presented) A DNzyme according to claim 2  
2    which includes an inverse thymidine on the 3' end and/or a FAM  
3    label on the 5' end.

1                   8. (previously presented) A medicament containing a  
2    DNzyme according to claim 2 and a pharmaceutically acceptable  
3    carrier.

Claims 9 through 16 (canceled).